

K812688 QI 1150-1T INFUSION TOct 2, 1981
10 days to decisionK812688 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k812688/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Sep 22, 1981
Decision date	Oct 2, 1981
Days to decision	10 days
Third-party review	No

APPLICANT

Company	Quinton, Inc.
Location	Mchenry, IL, US
510(k) history	164 submissions · 160 cleared · 1976-2003

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Device record: <https://www.510kdatabase.net/k812688/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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